Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID: SSSPTA1623PAZ

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

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Welcome to STN International
NEWS
                 Web Page URLs for STN Seminar Schedule - N. America
      1
NEWS
                 "Ask CAS" for self-help around the clock
                BEILSTEIN enhanced with new display and select options,
NEWS
                 resulting in a closer connection to BABS
                IFIPAT/IFIUDB/IFICDB reloaded with new search and display
NEWS
        AUG 02
                 fields
NEWS
     5
        AUG 02
                 CAplus and CA patent records enhanced with European and Japan
                 Patent Office Classifications
NEWS
     6
        AUG 02
                 The Analysis Edition of STN Express with Discover!
                 (Version 7.01 for Windows) now available
                 BIOCOMMERCE: Changes and enhancements to content coverage
NEWS
     7
         AUG 27
                BIOTECHABS/BIOTECHDS: Two new display fields added for legal
NEWS
         AUG 27
                 status data from INPADOC
                 INPADOC: New family current-awareness alert (SDI) available
NEWS 9
         SEP 01
                 New pricing for the Save Answers for SciFinder Wizard within
NEWS 10
        SEP 01
                 STN Express with Discover!
                 New display format, HITSTR, available in WPIDS/WPINDEX/WPIX
NEWS 11
        SEP 01
NEWS 12
         SEP 27
                 STANDARDS will no longer be available on STN
                 SWETSCAN will no longer be available on STN
NEWS 13
         SEP 27
NEWS 14 OCT 28
                 KOREAPAT now available on STN
             OCTOBER 29 CURRENT WINDOWS VERSION IS V7.01A, CURRENT
NEWS EXPRESS
              MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 11 AUGUST 2004
NEWS HOURS
              STN Operating Hours Plus Help Desk Availability
              General Internet Information
NEWS INTER
NEWS LOGIN
              Welcome Banner and News Items
NEWS PHONE
              Direct Dial and Telecommunication Network Access to STN
NEWS WWW
              CAS World Wide Web Site (general information)
```

Enter NEWS followed by the item number or name to see news on that specific topic.

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FILE 'HOME' ENTERED AT 07:24:52 ON 10 NOV 2004

=> file reg
COST IN U.S. DOLLARS
SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST
0.21
0.21

FILE 'REGISTRY' ENTERED AT 07:25:39 ON 10 NOV 2004

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9 DICTIONARY FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

=> logoff hold COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 2.94 3.15

FULL ESTIMATED COST

SESSION WILL BE HELD FOR 60 MINUTES
STN INTERNATIONAL SESSION SUSPENDED AT 07:29:37 ON 10 NOV 2004

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID:SSSPTA1623PAZ

PASSWORD:

* * * * * RECONNECTED TO STN INTERNATIONAL * * * * * * * * * SESSION RESUMED IN FILE 'REGISTRY' AT 07:31:58 ON 10 NOV 2004 FILE 'REGISTRY' ENTERED AT 07:31:58 ON 10 NOV 2004 COPYRIGHT (C) 2004 American Chemical Society (ACS)

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 2.94 3.15

FULL ESTIMATED COST

Uploading C:\Examination Auxillary files\10815375\10815375 core structure.str

chain nodes :

1 2 3 4 5 6 7 8 9 10 11 12

chain bonds :

1-2 2-3 2-10 3-4 4-5 5-6 6-7 7-8 8-9 8-11 8-12

exact/norm bonds :

2-3 2-10 3-4 5-6 6-7 8-12

exact bonds :

1-2 4-5 7-8 8-9 8-11

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS 11:CLASS 12:CLASS

L1 STRUCTURE UPLOADED

=> d l;1

L1 HAS NO ANSWERS

'L ' IS NOT A VALID STRUCTURE FORMAT KEYWORD

Structure Formats

SIA ---- Structure Image, Attributes, and map table if it contains data. (Default)

SIM ---- Structure IMage.

SAT ---- Structure ATtributes and map table if it contains data.

SCT ---- Structure Connection Table and map table if it contains data.

SDA ---- All Structure DAta (image, attributes, connection table and map table if it contains data).

NOS ---- NO Structure data.

ENTER STRUCTURE FORMAT (SIM), NOS:end

SYSTEM LIMITS EXCEEDED - SEARCH ENDED

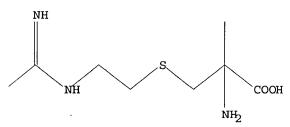
The search profile you entered was too complex or gave too many answers. Simplify or subdivide the query and try again. If you have exceeded the answer limit, enter DELETE HISTORY at an arrow prompt (=>) to remove all previous answers sets and begin at L1. Use the SAVE command to store any important profiles or answer sets before using DELETE HISTORY.

=> d 11

L1 HAS NO ANSWERS

Ll

STR



Structure attributes must be viewed using STN Express query preparation.

=> search l1 sss sam

SAMPLE SEARCH INITIATED 07:32:43 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 12 TO ITERATE

100.0% PROCESSED

12 ITERATIONS

9 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 33 TO 447 PROJECTED ANSWERS: 9 TO 360

=> d scan

9 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN L2

L-Cysteine, S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-, carbonate (1:1) IN

C8 H17 N3 O2 S . C H2 O3 MF

> CM 1

Absolute stereochemistry.

$$\begin{array}{c|c} NH & Me \\ NH_2 & NH_2 \\ Me & NH_2 \\ \\ Me & CO_2H \end{array}$$

CM 2

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):10

REGISTRY COPYRIGHT 2004 ACS on STN 9 ANSWERS L2

L-Cysteine, S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-, IN 2-hydroxy-1,2,3-propanetricarboxylate (1:1) (9CI)

MF C8 H17 N3 O2 S . C6 H8 O7

> CM 1

Absolute stereochemistry.

$$\begin{array}{c|c} NH & \text{Me} \\ \hline NH_2 & \\ NH_2 & \\ CO_2H & \\ \end{array}$$

CM 2

$$\begin{array}{c} {\rm CO_2H} \\ | \\ {\rm HO_2C-CH_2-C-CH_2-CO_2H} \\ | \\ {\rm OH} \end{array}$$

REGISTRY COPYRIGHT 2004 ACS on STN ·L2 9 ANSWERS L-Cysteine, S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-, IN (2S,3S)-2,3-dihydroxybutanedioate (1:1) (9CI) C8 H17 N3 O2 S . C4 H6 O6 MF

Absolute stereochemistry.

$$\begin{array}{c|c} NH & Me \\ NH_2 \\ CO_2H \end{array}$$

CM 2

Absolute stereochemistry.

L29 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

IN D-Isovaline, 2'-[[2-[(1-iminoethyl)amino]ethyl]thio]-3-methyl-,

dihydrochloride (9CI)

MF C10 H21 N3 O2 S . 2 Cl H

Absolute stereochemistry.

●2 HCl

9 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN L-Cysteine, S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-, compd. with IN

L-ascorbic acid (1:1) (9CI)

C8 H17 N3 O2 S . C6 H8 O6 MF

> CM1

Absolute stereochemistry.

Absolute stereochemistry.

9 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN L2

Cysteine, S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-, IN

bis(trifluoroacetate) (9CI) C8 H17 N3 O2 S . 2 C2 H F3 O2

CM

MF

$$\begin{array}{c|c} \mathrm{NH_2} & \mathrm{NH} \\ | & | \\ \mathrm{HO_2C-C-CH_2-S-CH_2-CH_2-NH-C-Me} \\ | & \\ \mathrm{Me} \end{array}$$

CM

REGISTRY COPYRIGHT 2004 ACS on STN L2

9 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN L-Cysteine, S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-, IN

(2R)-hydroxybutanedioate (20:11) (9CI)

C8 H17 N3 O2 S . 11/20 C4 H6 O5 MF

> CM 1

Absolute stereochemistry.

CM 2

Absolute stereochemistry.

L2

9 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN L-Cysteine, S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-, IN

mono(3-phosphonopropanoate) (9CI)

C8 H17 N3 O2 S . C3 H7 O5 P ΜF

> CM 1

Absolute stereochemistry.

Me NH₂
$$\sim$$
 NH₂ \sim CO₂H

CM 2

HO2C-CH2-CH2-PO3H2

REGISTRY COPYRIGHT 2004 ACS on STN L2

L-Cysteine, S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-, monohydrochloride IN mono[(2S)-2-hydroxypropanoate] (9CI)

C8 H17 N3 O2 S . C3 H6 O3 . C1 H MF

> CM 1

Absolute stereochemistry.

CM 2

Absolute stereochemistry. Rotation (+).

ALL ANSWERS HAVE BEEN SCANNED

=> search l1 sss full FULL SEARCH INITIATED 07:34:17 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 181 TO ITERATE 100.0% PROCESSED 181 ITERATIONS 164 ANSWERS

SEARCH TIME: 00.00.01

164 SEA SSS FUL L1 L3

=> file caplus COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 159.62 159.83

FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 07:34:26 ON 10 NOV 2004 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2004 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 10 Nov 2004 VOL 141 ISS 20 FILE LAST UPDATED: 8 Nov 2004 (20041108/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> 13

L415 L3

=> d l4 1-15 ti

- ANSWER 1 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- Exchanger for selectively removing counterions from compounds and compounds derived from the methods for pharmaceutical applications
- ANSWER 2 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-cysteine maleate hydrochloride ΤI crystalline salt
- ANSWER 3 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
- S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-cysteine maleate form II ΤI crystalline salt
- ANSWER 4 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-cysteine maleate form II TI crystalline salt
- ANSWER 5 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-cysteine salicylate TTmonohydrate crystalline salt
- ANSWER 6 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- Solid dispersions comprising a hygroscopic and/or deliquescent drug TI
- ANSWER 7 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- Non-hygroscopic formulation comprising a hygroscopic drug TI

- ANSWER 8 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- Preparation of 2,7-diamino-5-heptenoic acid derivatives for the treatment TΙ and prevention of gastrointestinal conditions
- ANSWER 9 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN **L4**
- TI Preparation of amino acid derivatives and methods for the treatment of respiratory diseases and conditions using a selective inos inhibitor
- ANSWER 10 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN **L4**
- Preparation of amino acid derivatives in methods for the treatment of ΤI respiratory diseases and conditions with a selective iNOS inhibitor and a PDE inhibitor
- ANSWER 11 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- Preparation of amino acid derivatives as selective nitric oxide synthase TIinhibitors for ophthalmol. treatment
- ANSWER 12 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- Preparation of amino acid derivatives as selective nitric oxide synthase ΤI inhibitors for neuroprotective treatment
- L4ANSWER 13 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
- TIPreparation of 2,7-diamino-5-heptenoic acid derivatives for the treatment of cancer
- ANSWER 14 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- Preparation of amidino compounds useful as nitric oxide synthase TTinhibitors
- ANSWER 15 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN L4
- ΤI Preparation of amidino compounds useful as nitric oxide synthase inhibitors

=> d 14 12-15 ti fbib abs

- ANSWER 12 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
- Preparation of amino acid derivatives as selective nitric oxide synthase ΤI inhibitors for neuroprotective treatment
- AN 2003:261658 CAPLUS
- 138:287965 DN
- Preparation of amino acid derivatives as selective nitric oxide synthase TI inhibitors for neuroprotective treatment
- Manning, Pamela T.; Connor, Jane R. IN
- Pharmacia Corporation, USA PA
- PCT Int. Appl., 185 pp. SO CODEN: PIXXD2
- DT Patent
- LΑ English

F	AN.CNT	1																
	PA	TENT	NO.			KIN	D	DATE		1	APPL	ICAT	ION 1	NO.		D	ATE	
				~			_		-							_		
P	I WO	2003	0266	38		A1		2003	0403	1	WO 2	002-1	US30:	214		2	0020	924
		W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,
			CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	LK,	LR,
			LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,
			PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TN,	TR,	TT,	TZ,
			UA,	UG,	US,	UZ,	VN,	YU,	ZA,	ZM,	ZW							
		RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	AZ,	BY,
			KG,	KΖ,	MD,	RU,	TJ,	TM,	ΑT,	ΒE,	CH,	CY,	DE,	DK,	ES,	FI,	FR,	GB,
			GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,
			GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	TG							
										1	US 2	001-	96152	21	Ī	A 20	00109	924
	ITC	2002	1100	26		7. 1		2002	0606	1	TC 2	001	0615	2.1		2.	2010	224

EP 1429752 A1 20040623 EP 2002-761804 20020924 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

US 2001-961521 A 20010924 WO 2002-US30214 W 20020924

OS MARPAT 138:287965

GΙ

The acetamidino amino acid derivs. I [R5 = H, or OH; X = CR1:CR2CH2, CR1:CR2, CH2CR1:CR2, C.tplbond.C, CH2C.tplbond.C, C.tplbond.CCH2; R1, R2 = H, halo, alkyl, or haloalkyl; with the proviso that at least one of R1 or R2 contains halo; Z = H, (un) substituted alkyl, alkoxy, or halo; J = H, OH, alkoxy, NR3R4; R3 = H, alkyl, alkenyl, alkynyl; R4 = H, or (un) substituted heterocyclyl] and related 7-iminohexahydro-2-azepinyl derivs. were prepared as selective nitric oxide synthase inhibitors for the prevention and treatment of neurodegenerative conditions. Thus, (2S,5E)-2-amino-6-fluoro-7-[(1-iminoethyl)amino]-5-heptenoic acid dihydrochloride prepared by a multistep procedure starting from L-glutamic acid inhibited the LPS-induced increase in plasma nitrite/nitrate levels with an observed ED50 value of <0.1 mg/kg demonstrating the ability to inhibit inducible nitric oxide synthase activity in vivo.

RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 13 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN

TI Preparation of 2,7-diamino-5-heptenoic acid derivatives for the treatment of cancer

AN 2002:754159 CAPLUS

DN 137:263297

TI Preparation of 2,7-diamino-5-heptenoic acid derivatives for the treatment of cancer

IN Manning, Pamela T.; Connor, Jane R.; Seibert, Karen; Rao, Chinthalapally
 V.; Reddy, Bandaru S.

PA Pharmacia Corporation, USA

SO PCT Int. Appl., 295 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT	NO.			KIN	D :	DATE			APPL	ICAT:	ION I	MO.		D	ATE	
						-									-		
ΡI	WO 2002	07639	95		A2		2002	1003	1	WO 2	002-1	US89:	38		2	0020	321
	WO 2002	0763	95		A 3		2004	0812									
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		CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KΡ,	KR,	KZ,	LC,	LK,	LR,
		LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,
		PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,	TZ,
		UA,	UG,	US,	UΖ,	VN,	YU,	ZA,	ZM,	ZW,	AM,	ΑZ,	BY,	KG,	KZ,	MD,	RU,
,		ТJ,	TM														
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		BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG
									1	US 20	001-2	2785	12P	I	2 (00103	323
									1	US 20	001-9	96196	59	1	A 20	00109	924
	US 2003	01370)2		Al	:	2003	0116	1	US 20	001-	96196	59		20	00109	924

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P 20010323
                                       US 2001-278512P
                           20041006
EP 1463495
                     A2
                                       EP 2002-717708
                                                              20020321
    R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
        IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                                       US 2001-278512P
                                                           P
                                                              20010323
                                                              20010924
                                       US 2001-961969
                                                           Α
                                       WO 2002-US8938
                                                              20020321
```

OS MARPAT 137:263297

AB Agents and methods for chemoprevention and treatment of neoplasia are described, the agents including a selective inhibitor of inducible nitric oxide synthase and a combination of a selective inhibitor of inducible nitric oxide synthase and an inhibitor of cyclooxygenase-2 in a pharmaceutical composition 2,7-Diamino-5-heptenoic acid derivs.

R7N:CMeNHCH2CR1:CR2CH2CH2CH(NH2)C(O)J [R1, R2 = H, halo, alkyl, haloalkyl (at least one of R1 or R2 contains halogen); R7 = H, OH; J = OH, alkoxy, NR3R4, where R3 = H, alkyl, alkenyl, alkynyl and R4 = H, (un)substituted heterocyclyl] or their pharmaceutically-acceptable salts are among the compds. claimed. Thus, (2S,5E)-2-amino-6-fluoro-7-[(1-iminoethyl)amino]-5-heptenoic acid dihydrochloride was prepared by a multistep procedure starting from L-glutamic acid and showed IC50 values 0.36, 68, 3.6, and 0.1 μM in hiNOS, hecNOS, hncNOS, and human cartilage assays, resp.

- L4 ANSWER 14 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Preparation of amidino compounds useful as nitric oxide synthase inhibitors
- AN 2001:730698 CAPLUS
- DN 135:289056
- TI Preparation of amidino compounds useful as nitric oxide synthase inhibitors
- IN Webber, Ronald Keith; Awasthi, Alok K.; Bergmanis, Arija A.; Durley, Richard C.; Ganser, Scott S.; Hagen, Timothy J.; Hallinan, Ann E.; Hansen, Donald W.; Hickory, Brian S.; Moormann, Alan E.; Pitzele, Barnett S.; Promo, Michelle A.; Schartman, Richard R.; Snyder, Jeffrey S.; Trivedi, Mahima; Tsymbalov, Sofya
- PA Pharmacia Corporation, USA
- SO PCT Int. Appl., 144 pp. CODEN: PIXXD2
- DT Patent
- LA English

FAN CNT 2

FAN.	CNT	2				•												
	PAT	CENT 1	NO.			KIN	D	DATE				ICAT				D	ATE	
ΡI	WO	2001	0727	03		A1										2	0010	323
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			CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,
			HR,	ΗU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	KP,	KR,	ΚZ,	LC,	LK,	LR,	LS,
			LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	PL,	PT,	RO,
			RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TR,	TT,	TZ,	UA,	UG,	US,	UΖ,
			VN,	YU,	ZA,	ZW,	AM,	ΑZ,	BY,	KG,	KZ,	MD,	RU,	ТJ,	TM			
		RW:	GH,	GM,	KΕ,	LS,	MW,	ΜZ,	SD,	SL,	SZ,	TZ,	UG,	ZW,	ΑT,	BE,	CH,	CY,
			DE,	DK,	ES,	FI,	FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	TR,	BF,
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										1	US 2	000-	19192	23P]	P 2	00003	324
	US	2002	0195	63		A1		2002	0214	1	US 2	001-	8165	77		2	0010	323
	US	6403	830			B2		2002	0611									
										1	US 2	000-	1919:	23P]	P 2	0000	324
	US	2002	1114	93		A1		2002	0815	1	US 2	001-	8165'	75		2	0010	323
	US	6586	474			B2		2003	0701									
										1	US 2	000-	1919:	23P]	P 2	0000	324
	EP	1265	860			A1		2002	1218	:	EP 2	001-	92263	36		2	0010	323
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			ΙE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	\mathbf{T} R						
										1	US 2	000-	19192	23P]	P 2	00003	324
										1	WO 2	001-1	US94:	33	1	V 2	0010	323
	BR	2001	0093	86		Α		2003	0415]	BR 2	001-	9386			2	00103	323

						US 2000-191923P P 20000324
						WO 2001-US9433 W 20010323
	ZA 20020	006459		Α	20030813	
	TD 20031			Т2	20030930	US 2000-191923P P 20000324 JP 2001-570616 20010323
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		HR, HU,	ID,	IL,	IN, IS, JP,	KE, KG, KP, KR, KZ, LC, LK, LR, LS,
		LT, LU,	LV,	MA,	MD, MG, MK,	MN, MW, MX, MZ, NO, NZ, PL, PT, RO,
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	RW.					IE, IT, LU, MC, NL, PT, SE, TR, BF,
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						US 2002-321969 B3 20021217
os	MARPAT :	135:2890:	56			

OS MARPAT 135:289056 AB

The invention relates to S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-

cysteine (1) or its pharmaceutically acceptable salts for use as nitric oxide synthase (NOS) inhibitors. Thus, 1.2HCl was prepared by a multistep procedure involving S-alkylation of (2R)-2-methyl-L-cysteine hydrochloride with Boc-NHCH2CH2Br (Boc = tert-butoxycarbonyl), deprotection, condensation with Et acetimidate hydrochloride, and acidolysis with 1 N HCl. (2R)-2-methyl-L-cysteine hydrochloride was obtained from (R)-cysteine Me ester hydrochloride. Inhibitory assays for compound 1.2HCl showed hiNOS, hecNOS, hncNOS, and human cartilage IC50 values 3.1, 77, 15 μM , and 0.7 μM , resp.

RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L4 ANSWER 15 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Preparation of amidino compounds useful as nitric oxide synthase inhibitors
- AN 2001:730697 CAPLUS
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- TI Preparation of amidino compounds useful as nitric oxide synthase inhibitors
- IN Webber, Ronald Keith; Awasthi, Alok K.; Bergmanis, Arija A.; Durley, Richard C.; Fok, Kam F.; Ganser, Scott S.; Hagen, Timothy J.; Hallinan, Ann E.; Hansen, Donald W.; Hickory, Brian S.; Manning, Pamela T.; Mao, Michael; Moormann, Alan E.; Pitzele, Barnett S.; Promo, Michelle A.; Schartman, Richard R.; Scholten, Jeffrey A.; Snyder, Jeffrey S.; Toth, Mihaly V.; Trivedi, Mahima; Tsymbalov, Sofya; Tjoeng, Foe Siong
- PA Pharmacia Corporation, USA
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- LA English

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	NZ 520	813	Ĭ	Ą	2004	0528	I J	NZ 2 US 2	001- 000-	5208 1919	13 23P		2 P 2	0010 0000 0010	323 324
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Amidino compds. R11N:CR13NR12CR9R10CR1R7-X-CR5R6CR2(NR3R4)COR8 [X = S, SO, SO2; R1, R5, R6, R7 = H, halo, alkyl (alkyl and other groups may be substituted), alkenyl, alkynyl, alkoxyalkyl; R2 = alkyl, alkenyl, alkynyl, alkoxyalkyl, alkylthioalkyl; R3 = H, OH, CHO, alkanoyl, CO2H, C(O)SH or alkyl esters; R8 = OH, alkoxy, an amino or alkylamino group or R3 and R8 may form a ring; R4 = H, CO2H, carbalkoxy; R9, R10 = H, alkyl, alkenyl, alkynyl, alkoxyalkyl; R11, R12 = H, OH, CO2H, C(O)SH or esters or R11 and R12 may form a ring; R13 = alkyl (with provisos)] or their salts were prepared as nitric oxide synthase (NOS) inhibitors. Thus, S-[2-[(1-iminoethyl)amino]ethyl]-2-methyl-L-cysteine dihydrochloride (1)

was prepared by a multistep procedure involving S-alkylation of (2R)-2-methyl-L-cysteine hydrochloride with Boc-NHCH2CH2Br (Boc = tert-butoxycarbonyl), deprotection, condensation with Et acetimidate hydrochloride, and acidolysis with 1 N HCl. (2R)-2-methyl-L-cysteine hydrochloride was obtained from (R)-cysteine Me ester hydrochloride. Inhibitory assays for compound 1 showed hiNOS, hecNOS, hncNOS, and human cartilage IC50 values 3.1, 77, 15 μM , and 0.7 μM , resp.

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SESSION WILL BE HELD FOR 60 MINUTES
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13	BRS	L13	0	"9962875".pn.	USPAT	2004/11/10 07:56	
14	BRS	L37	248	564/197.ccls.	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
15	BRS	L38	843	514/562.ccls.	USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
16	BRS	L39	311636	beams	USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
17	BRS	L40	11	nitic adj oxide	USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
18	BRS	L41	10811	nitric adj oxide	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
19	BRS	L42	6072	amidino	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	

20	BRS	L43	1151	amidino and (nitric adj oxide)	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
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	Туре	L #	Hits	Search Text	DBs	Time Stamp	Comments
21	BRS	L44	138	548/316.7	USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
22	BRS	L45	259	560/153.ccls.	USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
23	BRS	L46	432	560/147.ccls.		2004/11/10 07:56	
24	BRS	L47	136	560/149.ccls.	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
25	BRS	L48	911	548/316.7 or 560/153.ccls. or 560/147.ccls. or 560/149.ccls.	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
26	BRS	L49	30345	cysteine	USPAT	2004/11/10 07:56	
27	BRS	L50	o	"9962875".pn.	USPAT	2004/11/10 07:56	
28	BRS	L51	o	"9962875".pn.	USPAT	2004/11/10 07:56	
29	BRS	L52	248	564/197.ccls.	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
30	BRS	L53	843	514/562.ccls.	USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
31	BRS	L18	1	"5863931".PN.	USPAT	2004/11/10 07:56	
32	BRS	L19	1	"5684008".PN.	USPAT	2004/11/10 07:56	

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	Туре	L #	Hits	Search Text	DBs	Time Stamp	Comments
33	BRS	L21	1	"9324126".pn.	LTD() •	2004/11/10 07:56	
34	BRS	L23	1	"5854234".pn.		2004/11/10 07:56	
35	BRS	L25	1	"6046211" .pn.		2004/11/10 07:56	
36	BRS	L26	1	"5674894".pn.	ILI Q D A Tr	2004/11/10 07:56	
37	BRS	L32	1	"5863931".PN.	יניאכרים	2004/11/10 07:56	
38	BRS	L33	1	"5889056".PN.	III CODATI	2004/11/10 07:56	
39	BRS	L34	1	"2049582".PN.	USPAT	2004/11/10 07:56	
40	BRS	L35	1	"4085217".PN.	USPAT	2004/11/10 07:56	
41	BRS	L54	1	"5863931".PN.	USPAT	2004/11/10 07:56	
42	BRS	L55	1	"5684008".PN.	USPAT	2004/11/10 07:56	
43	BRS	L56	1	"9324126".pn.	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
44	BRS	L57	1	"5854234".pn.	USPAT	2004/11/10 07:56	
45	BRS	L58	1	"6046211" .pn.	USPAT	2004/11/10 07:56	
46	BRS	L59	1	"5674894".pn.	USPAT	2004/11/10 07:56	
47	BRS	L60	1	"5863931".PN.	USPAT	2004/11/10 07:56	
48	BRS	L61	1	"5889056".PN.	USPAT	2004/11/10 07:56	
49	BRS	L62	1	"2049582".PN.	USPAT	2004/11/10 07:56	
50	BRS	L63	1	"4085217".PN.	USPAT	2004/11/10 07:56	

51	IS&R	L14	2	("6403830").PN.		2004/11/10 07:56	
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52	IS&R	L15	2	("6355689").PN.		2004/11/10 07:56	
53	BRS	L16	2	"9313055".pn.	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
54	BRS	L20	3	"9525717".pn.	110()	2004/11/10 07:56	
55	BRS	L22	39	(amidino and (nitric adjoxide)) and cysteine	USPAT	2004/11/10 07:56	
56	BRS	L24	12	(nitric adj oxide) and (548/316.7 or 560/153.ccls. or 560/147.ccls. or 560/149.ccls.)	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
57	BRS	L27	3	"9962875".pn.	US- PGPUB; USPAT; EPO; JPO; DERWEN	07:56	
58	IS&R	L28	2	("6355689").PN.	USPAT; USOCR; EPO; JPO; DERWEN	2004/11/10 07:56	

59	IS&R	L29	2	("6403830").PN.		2004/11/10 07:56	
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60	BRS	L30	3	"2000044731".pn.	IP. DU .	2004/11/10 07:56	
61	BRS	L31	9	(nitric adj oxide) and 562/557.ccls.	114 114 114	2004/11/10 07:56	
62	IS&R	L64	2	("6403830").PN.		2004/11/10 07:56	
63	IS&R	L65	2	("6355689").PN.	USPAT; USOCR; EPO; JPO; DERWEN T	2004/11/10 07:56	
64	BRS	L66	2	"9313055".pn.	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
65	BRS	L67	3	"9525717".pn.	USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
66	BRS	L68	39	(amidino and (nitric adjoxide)) and cysteine	USPAT	2004/11/10 07:56	
67	BRS	L 69	12	(nitric adj oxide) and (548/316.7 or 560/153.ccls. or 560/147.ccls.)	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	

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	Туре	L #	Hits	Search Text	DBs	Time Stamp	Comments
68	BRS	L70	3	, "9962875".pn.	16' D() -	2004/11/10 07:56	
69	IS&R	L71	2	("6355689").PN.		2004/11/10 07:56	
70	IS&R	L72	2	("6403830").PN.		2004/11/10 07:56	
71	BRS	L73	3	"2000044731".pn.	US- PGPUB; USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	,
72	BRS	L74	9	(nitric adj oxide) and 562/557.ccls.	US- PGPUB; USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
73	BRS	L77	5	"6586474"	US- PGPUB; USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	

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74	BRS	L78	2	"6586474".pn.	IN DO +	2004/11/10 07:56	
75	BRS	L17	186	beams and (nitric adj oxide)	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
76	BRS	L75	186	beams and (nitric adj oxide)	USPAT; EPO; JPO; DERWEN T	2004/11/10 07:56	
77	BRS	L36	241	562/557.ccls.	US- PGPUB; USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
78	BRS	L76	241	562/557.ccls.	US- PGPUB; USPAT; EPO; JPO; DERWEN	2004/11/10 07:56	
79	BRS	L 79	13	US-5132453-\$.DID. OR US- 5684008-\$.DID. OR US- 5830917-\$.DID. OR US- 5854251-\$.DID. OR US- 5863931-\$.DID. OR US- 5919787-\$.DID. OR US- 5945408-\$.DID. OR US- 5981511-\$.DID.		2004/11/10 08:35	

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